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PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Application No.:

10/735,408

Confirmation No.: 2099

Applicant:

Storer et al.

Filed:

December 12, 2003

TC/A.AU.: Examiner:

Unassigned Unassigned

Docket No.:

06171.105101 IDX 1024

Customer No.:

20786

Title:

Process for the Production of 2'-Branched Nucleosides

Commissioner for Patents

P. O. Box 1450

Alexandria, VA 22313-1450

Information Disclosure Statement

Sir:

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Applicants do not believe any fees are due because this paper is submitted before the mailing of a first Office action on the merits, as under 37 C.F.R. § 1.97(b)(3). However, the Commissioner is hereby authorized to charge any fees due or credit any overpayment to Deposit Account No. 11-0980.

Respectfully submitted,

of explos person Madely fluid 36,174 Sherry M. Knowles

Reg. No. 33,052

Dated: September 10, 2004 King & Spalding, LLP

191 Peachtree Street, N.E., Atlanta, GA 30303 Office: (404)572-4600/ Fax: 404-572-5145

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Brent R. Bellows

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Substitute for form 1449A/PTO

INFORMATION DISCLOSURE STATEMENT BY APPLICANT

(use as many sheets as necessary)

1 of 8

Complete if Known					
Application Number	10/735,408				
Filing Date	December 12, 2003				
First Named Inventor	Storer et al.				
Group Art Unit	1623				
Examiner Name	Unassigned				
Attorney Docket Number	06171.105101 IDX 1024				

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Examiner Initials *	Cite No. 1		nent Kind Code (if known)	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages/Relevant Figures Appear	T6
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¹ Unique citation designation number. ² See attached Kinds of U.S. Patent Documents. ³ Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). ⁴ For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. ⁵ Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. ⁶ Applicant is to place a check mark here if English language Translation is attached.

Complete if Known Substitute for form 1449A/PTO Application Number 10/735,408 Filing Date INFORMATION DISCLOSURE **December 12, 2003** First Named Inventor STATEMENT BY APPLICANT Storer et al. Group Art Unit 1623 (use as many sheets as necessary) **Examiner Name** Unassigned **Attorney Docket Number** 06171.105101 IDX 1024 8

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	CK	WO	04/009020	A2	Merck & Co., Isis Pharmaceutical	01-29-2004		

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Examiner Initials *	Cite No. 1	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ⁶
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Substitut	Substitute for form 1449A/PTO INFORMATION DISCLOSURE STATEMENT BY APPLICANT (use as many sheets as necessary)		Application Number	10/735,408	
INF	ORMATION I	DISCL	OSURE	Filing Date	December 12, 2003
STA	TEMENT BY	APPL	ICANT	First Named Inventor	Storer et al.
				Group Art Unit	1623
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		OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS	
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Examiner	Date	·
Signature	Considered	

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Substitute for form 1449A/PTO	NFORMATION DISCLOSURE TATEMENT BY APPLICANT (use as many sheets as necessary)		Application Number	10/735,408
INFORMATION	N DISCL	OSURE	Filing Date	December 12, 2003
STATEMENT B	Y APPL	ICANT	First Named Inventor	Storer et al.
			Group Art Unit	1623
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5	of	8	Attorney Docket Number	06171.105101 IDX 1024

		OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS	
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Examiner Signature	·	Date Considered	

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Substitute for form 1449A/PTO			Complete if Known		
Substitute for form 1449A/F1O			Application Number	10/735,408	
INFORMATION 1	DISCL	OSURE	Filing Date	December 12, 2003	
STATEMENT BY	APPL	ICANT	First Named Inventor	Storer et al.	
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		OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS	
Examiner Initials *	Cite No. 1	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ⁶
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¹ Unique citation designation number. ² See attached Kinds of U.S. Patent Documents. ³ Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). ⁴ For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. ⁵ Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. ⁶ Applicant is to place a check mark here if English language Translation is attached.

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STATEMENT B	STATEMENT BY APPLICANT			Storer et al.		
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7 of 8			Attorney Docket Number	06171.105101 IDX 1024		
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OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, Examiner Cite T⁶ serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published. Initials * No. 1 LOPEZ APARICIO, F.J., et al., "Synthesis of saccharinic acid derivatives," Carbohydrate Res., GA 129:99 (1984), LUH, T.-Y., et al., "A convenient method for the selective esterification of amino-alcohols," GB Synthetic Communications, 8(5):327-333 (1978). McCORMICK, J., et al., "Structure and total synthesis of HF-7, a neuroactive glyconucleoside GC disulfate from he funnel-web spide Hololena curta," J. Am. Chem. Soc., 121(24), 5661-5664 MCKENZIE, R., et al., "Hepatic failure and lactic acidosis due to fialuridine (FIAU), an GD investigational nucleoside analogue for chronic hepatitis B", N. Engl. J. Med., 333(17):1099-1105 (1995). MEDINA, D. J., et al., "Comparison of mitochondrial morphology, mitochondrial DNA content, **GE** and cell viability in cultured cells treated with three anti-Human Immunodeficiency Virus dideoxynucleosides," Antimicrob. Agents Chemother., 38(8):1824-8 (1994). MEIER, C., et al., "Cyclic saligenyl phosphotriesters of 2',3'-dideoxy-2',3'-didehydrothymidine GF (d4T) - A new pro-nucleic approach." Bioorganic & Med. Chem. Letters 7(2):99-104 (1997). MEYER, R.B., Jr., et al., "2'-O-Acyl-6-thioinosine cyclic 3',5'-phosphates as prodrugs of GG thioinosinic acid," J. Med. Chem. 22: 811-815 (1979). NEIDLEIN, R., et al., "Mild preparation of 1-benzyuloxyiminoalkylphosphonic dichlorides: GH Application to the synthesis of cyclic phosphonic diesters and cyclic monoester amides," Heterocycles 35:1185-1203 (1993). NOVÁK, J.J.K. & SORM, F., "Nucleic acid components and their analogues. CXX. 2-C-methyl-GI D-ribose and tis derivatives," Collection Czechoslov. Chem. Commun., 34:857-866 (1969). NOVÁK, J.J.K., "Chiroptical properties of 2-methyl-1,4-lactones; revised absolute configuration of 2-deoxy-2-C-methyl-erythro-D-pentono-1,4-lactones," Collection Czechoslov. Chem. Commun., 39:869-882 (1974). NUTT, R.F., et al., "Branched-chain sugar nucleosides. III. 3'-C-methyladenine", J.Org. Chem., GK 33:1789-1795 (1968). OLSEN, et al. (Oral Session V, Hepatitis C Virus, Flaviviridae; 16th International Conference on GL Antiviral Research (April 27, 2003, Savannah, Ga.) p A76). PAN-ZHOU, X-R, et al., "Differential effects of antiretroviral nucleoside analogs on **GM** mitochondrial function in HepG2 cells," Antimicrob. Agents Chemother. 44:496-503 (2000). PIANTADOSI, C., et al., "Synthesis and evaluation of novel ether lipid nucleoside conjugates GN for anti-HIV-1 activity, " J. Med. Chem. 34:1408-1414 (1991). PIERRA, C., et al., "Comparative studies of selected potential prodrugs of β-L-dC, a potent and selective anti-HBV agent," Antiviral Res., 50:A79 (2001), Abstract no. 138.

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